## Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (currently amended) A method for treating a patient having an immune dysfunction, said method comprising the steps of:
- (a) treating peripheral blood mononuclear cells with an effective amount of an aziridino-containing compound; and
  - (b) administering said peripheral blood mononuclear cells to said patient, thereby treating said immune dysfunction in said patient, and

wherein said immune dysfunction is cutaneous T-cell lymphoma, graft versus host disease, allograft rejection following organ transplantation, systemic lupus erythematosus, systemic sclerosis, inflammatory bowel disease, or rheumatoid arthritis.

Claim 2 (cancelled).

3. (original) The method of claim 1 wherein said compound has the formula (II):

$$R_4 \longrightarrow R_1 \longrightarrow R_1 \longrightarrow R_2 \longrightarrow R_5 \longrightarrow R_6 \longrightarrow R_1 \longrightarrow R_1 \longrightarrow R_1 \longrightarrow R_2 \longrightarrow R_2 \longrightarrow R_2 \longrightarrow R_3 \longrightarrow R_4 \longrightarrow R_1 \longrightarrow R_2 \longrightarrow R_2 \longrightarrow R_2 \longrightarrow R_3 \longrightarrow R_4 \longrightarrow R_1 \longrightarrow R_2 \longrightarrow R_2 \longrightarrow R_3 \longrightarrow R_4 \longrightarrow R_2 \longrightarrow R_4 \longrightarrow R_4 \longrightarrow R_4 \longrightarrow R_5 \longrightarrow R_4 \longrightarrow R_4 \longrightarrow R_5 \longrightarrow R_5 \longrightarrow R_6 \longrightarrow R_4 \longrightarrow R_5 \longrightarrow R_6 \longrightarrow R_1 \longrightarrow R_2 \longrightarrow R_2 \longrightarrow R_4 \longrightarrow R_5 \longrightarrow R_5$$

wherein each  $R_1$  is a divalent hydrocarbon moiety containing between 2 and 4 carbon atoms, inclusive; each of  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ , and  $R_6$  is, independently, H or a monovalent hydrocarbon moiety containing between 1 arid 4 carbon atoms; and n is an integer between 1 and 10, inclusive.

- 4. (original) The method of claim 1, wherein said compound is ethyleneimine dimer.
- 5. (original) The method of claim 1, wherein said compound is an ethyleneimine trimer.
- 6. (original) The method of claim 1, wherein said compound is an ethyleneimine tetramer.
- 7. (currently amended) The method of claim 1, wherein said compound has the formula (III):

$$R_{5} = \begin{bmatrix} R_{2} \\ R_{1} - N^{4} - H \cdot n - W \end{bmatrix}$$

$$R_{6} = \begin{bmatrix} R_{2} \\ R_{3} \end{bmatrix}_{n}$$

$$(III)$$

wherein each  $R_1$  is a divalent hydrocarbon moiety containing between 2 and 4 carbon atoms, inclusive; each of  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $B_6$   $R_6$ , and  $R_7$  is, independently, H or a monovalent hydrocarbon moiety containing between 1 and 4 carbon atoms; X is C1 or Br, Y is a

pharmaceutically acceptable counter anion; W is valency of Y; and n is an integer between 1 and 10, inclusive.

## 8. (cancelled)

- 9. (currently amended) A method for treating a patient having an immune dysfunction, said method comprising the steps of:
- (a) extracorporeally treating peripheral blood mononuclear cells from said patient with an effective amount of an aziridino-containing compound;
- (b) separately separating said peripheral blood mononuclear cells from said aziridino-containing compound; and
  - (c) administering said peripheral blood mononuclear cells to said patient, thereby treating said immune dysfunction in said patient, and

wherein said immune dysfunction is cutaneous T-cell lymphoma, graft versus host disease, allograft rejection following organ transplantation, systemic lupus erythematosus, systemic sclerosis, inflammatory bowel disease, or rheumatoid arthritis.

- 10. (original) A method for preventing inhibiting tissue transplantation or blood transfusion-associated graft-versus-host (GVH) disease in a patient, the method comprising the steps of:
- (a) extracorporeally treating a blood composition with an effective amount of an aziridino-containing compound; and
  - (b) administering said treated blood cell population to said patient,

thereby preventing inhibiting tissue transplantation or blood transfusion-associated GVH disease in said patient.

- 11. (original) The method of claim 10, wherein said blood composition comprises peripheral blood mononuclear cells (PBMC).
- 12. (original) The method of claim 10, wherein said blood composition is a non-leukoreduced blood cell concentrate.
- 13. (original) The method of claim 10, wherein said blood composition is a heterologous blood cell population.
- 14. (original) The method of claim 10, wherein said method further separating said aziridino-containing compound from said treated blood cell composition prior to administering said treated blood composition to said patient.
- 15. (original) The method of claim 14, wherein at least 99% of said aziridino-containing compound is removed from said treated blood cell composition prior to administering said treated blood composition to said patient.
  - 16. (original) The method of claim 10, wherein said compound has the formula (II):

$$\begin{array}{c|c}
R_4 & & \\
R_5 & & \\
R_6 & & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 \\
R_1 - N - \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_1 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_1 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_1 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
N & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
R_4 & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
R_3 & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
R_4 & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
R_3 & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
R_4 & \\
\end{array}$$

$$\begin{array}{c|c}
R_2 & \\
R_3 & \\
\end{array}$$

$$\begin{array}{c|c}
R_3 & \\
R_4 & \\
\end{array}$$

$$\begin{array}{c|c}
R_4 & \\
R_5 & \\
\end{array}$$

$$\begin{array}{c|c}
R_5 & \\
R_5 & \\
\end{array}$$

$$\begin{array}{c|c}
R_5 & \\
\end{array}$$

wherein each  $R_1$  is a divalent hydrocarbon moiety containing between 2 and 4 carbon atoms, inclusive; each of  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ , and  $R_6$  is, independently, H or a monovalent hydrocarbon moiety containing between 1 arid 4 carbon atoms; and n is an integer between 1 and 10, inclusive.

- 17. (original) The method of claim 10, wherein said compound is an ethyleneimine dimer.
- 18. (original) The method of claim 10, wherein said compound is an ethyleneimine trimer.
- 19. (original) The method of claim 10, wherein said compound is an ethyleneimine tetramer.
  - 20. (original) The method of claim 10, wherein said compound has the formula (III):

$$\begin{array}{c|c}
R_{5} & & \\
R_{1} & & \\
R_{1} & & \\
R_{3} & & \\
\end{array}$$

$$\begin{array}{c|c}
R_{2} \\
H & n/-W[Y^{W}] \\
\end{array}$$
(III)

wherein each  $R_1$  is a divalent hydrocarbon moiety containing between 2 and 4 carbon atoms, inclusive; each of  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $B_6$ , and  $R_7$  is, independently, H or a monovalent hydrocarbon moiety containing between 1 and 4 carbon atoms; X is C1 or Br, Y is a pharmaceutically acceptable counter anion; W is valency of Y; and n is an integer between 1 and 10, inclusive.

- 21. (cancelled)
- 22. (original) The method of claim 10, wherein said patient is a human.
- 23. (original) The method of claim 10, wherein said patient suffers from or is at risk for immune dysfunction.
- 24. (original) The method of claim 22, wherein said human patient suffers from or is at risk for immune dysfunction.
- 25. (currently amended) A method for preventing- treating graft-versus-host (GVH) disease in a patient, the method comprising the steps of:

- (a) treating a heterologous blood composition with an effective amount of an ethylene oligomer compound;
  - (b) removing said ethylene oligomer from said heterologous treated blood composition; and
  - (c) administering said treated blood cell population to said patient, thereby preventing treating GVH disease in said patient.
  - 26. (original) The method of claim 25, wherein said patient is a human.
- 27. (original) The method of claim 25, wherein said compound is an ethyleneimine dimer.
- 28. (original) The method of claim 25, wherein said compound is an ethyleneimine trimer.
- 29. (original) The method of claim 25, wherein said compound is an ethyleneimine tetramer.
- 30. (original) A method for treating graft-versus-host (GVH) disease in a patient, the method comprising the steps of:
- (a) treating a heterologous blood composition with an effective amount of an aziridino-containing compound; and
  - (b) administering said treated blood cell population to said patient,

thereby treating GVH disease in said patient.

- 31. (original) A method for preventing inhibiting tissue transplantation or blood transfusion-associated graft-versus-host (GVH) disease in a patient, the method comprising the steps of:
- (a) treating a heterologous blood composition with an effective amount of an ethylene oligomer compound;
  - (b) removing said ethylene oligomer from said heterologous treated blood composition; and
- (c) administering said treated blood cell population to said patient,

  thereby preventing inhibiting blood transfusion-associated or blood transfusionassociated GVH disease in said patient.
- 32. (original) A method for preventing an inhibiting a tissue transplantation or blood transfusion-associated alloantibody response in a patient, the method comprising the steps of:
- (a) treating a heterologous blood composition with an effective amount of an aziridino-containing compound; and
- (b) administering said treated blood cell population to said patient,
  thereby preventing inhibiting said blood transfusion-associated alloantibody response in said patient.

Claims 33-35 (cancelled)

Claim 36 (new) The method of claim 1, wherein the peripheral blood mononuclear cells are contacted with a non-viricidal amount of said aziridino-containing compound.